IN THE CLAIMS:

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Cancel claims 1-28 and 31-48. This leaves pending claims 29-30 and 49-69 as shown in the full listing of claims below.

Claims 1-28 (canceled)

Claim 29 (original): A layered tablet for the treatment of herpes simplex and conditions giving rise thereto, said layered tablet comprising an immediate-release layer and a sustained-release layer, and comprising the following as active ingredients distributed between said immediate-release layer and said sustained-release layer in the following approximate proportions expressed as relative weight percents:

	Immediate-Release Layer	Sustained-Release Layer
Magnesium L-ascorbate	40-60%	balance
2-Amino-2-deoxy-D-glucose	40-60%	balance
L-lysine monohydrochloride	40-60%	balance
N-acetyl-L-cysteine	40-60%	balance
Quercetin	40-60%	balance
L-Selenomethionine	100%	
Copper sulfate	100%	
Zinc picolinate	40-60%	balance

Claim 30 (original): A layered tablet for use as an oral dosage form, said layered tablet comprising an immediate-release layer and a sustained-release layer, and comprising the following as active ingredients distributed between said immediate-release layer and said sustained-release layer in the following approximate proportions expressed as relative weight percents:

	Immediate-Release Layer	Sustained-Release Layer
Magnesium taurate	40-60%	balance

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L-selenomethionine	100%	
2-Amino-2-deoxy-D-glucose	40-60%	balance
L-Lysine ascorbate	50-60%	balance
Copper sulfate	100%	u
Zinc lysinate	40-60%	balance
N-acetyl-L-cysteine	40-60%	balance
Quercetin	40-60%	balance

6 Claims 31-48 (canceled)

- Claim 49 (original): A unit dosage form for the treatment of herpes simplex
 and conditions giving rise thereto, said unit dosage form being a vaginal dosage form
 selected from the group consisting of vaginally appropriate suppositories, creams, tablets and
 gels, comprising as active ingredients:
 - (a) about 1.3 μg/mL to about 30 μg/mL of ascorbic acid,
- 12 (b) about 0.01 mg/mL to about 0.2 mg/mL of 2-amino-2-deoxy-D-glucose,
- 13 (c) about 0.06 μ g/mL to about 8.5 μ g/mL of zinc sulfate, and
- 14 (d) about 1.6 μg/mL to about 23 μg/mL of L-lysine hydrochloride.
- Claim 50 (original): A unit dosage form in accordance with claim 49 further comprising as an active ingredient copper sulfate in a concentration ranging from about 0.4 µg/mL to about 15 µg/mL.
- 1 Claim 51 (original): A unit dosage form in accordance with claim 49 further
 2 comprising as an active ingredient quercetin in a concentration ranging from about 0.12
- 3 μ g/mL to about 2.75 μ g/mL.

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1	Claim 52 (original): A unit dosage form in accordance with claim 49 further
2	comprising as an active ingredient heparin sodium in a concentration ranging from about 0.6
3	unit/mL to about 8 units/mL.
1	Claim 53 (original): A unit dosage form in accordance with claim 49 further
2	comprising as an active ingredient quercetin in a concentration ranging from about 0.12
3	μg/mL to about 2.75 μg/mL and heparin sodium in a concentration ranging from about 0.6
4	unit/mL to about 8 units/mL.
1	Claim 54 (original): A unit dosage form in accordance with claim 49 further
2	comprising as an active ingredient quercetin in a concentration ranging from about 0.12
3	μg/mL to about 2.75 μg/mL, heparin sodium in a concentration ranging from about 0.6
4	unit/mL to about 8 units/mL, and N-acetylcysteine in a concentration ranging from about 0.6
5	units/mL to about 8 units/mL.
1	Claim 55 (original): A unit dosage form for the treatment of herpes simplex
2	and conditions giving rise thereto, said unit dosage form being a mucosal dosage form
3	selected from the group consisting of vaginally appropriate suppositories, creams, tablets and
4	gels, comprising as active ingredients:
5	(a) ascorbic acid,
6	(b) 2-amino-2-deoxy-D-glucose,
7	(c) zinc sulfate, and
8	(d) L-lysine hydrochloride.
1	Claim 56 (original): A unit dosage form in accordance with claim 55 in which
2	the concentrations of said active ingredients are as follows:
3	(a) about 1.3 μg/mL to about 30 μg/mL of ascorbic acid,
4	(b) about 0.01 mg/mL to about 0.2 mg/mL of 2-amino-2-deoxy-D-glucose,

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5	(c) about 0.06 μg/mL to about 8.5 μg/mL of zinc sulfate, and
6	(d) about 1.6 μ g/mL to about 23 μ g/mL of L-lysine hydrochloride.
1	Claim 57 (original): A unit dosage form in accordance with claim 55 further
2	comprising as an active ingredient copper sulfate in a concentration ranging from about 0.4
3	μg/mL to about 15 μg/mL.
1	Claim 58 (original): A unit dosage form in accordance with claim 55 further
2	comprising as an active ingredient heparin sodium in a concentration ranging from about 0.6
3	units/mL to about 8 units/mL.
1	Claim 59 (original): A unit dosage form in accordance with claim 55 further
2	comprising as active ingredients copper sulfate in a concentration ranging from about 0.4
3	μg/mL to about 15 μg/mL and heparin sodium in a concentration ranging from about 0.6
4	units/mL to about 8 units/mL.
1	Claim 60 (original): A unit dosage form in accordance with claim 55 further
2	comprising as an active ingredient N-acetyl-L-cysteine in a concentration ranging from about
3	0.02 mg/mL to about 0.5 mg/mL.
1	Claim 61 (original): A unit dosage form in accordance with claim 55 further
2	comprising as an active ingredient L-2-oxothiazolidine-4-carboxylate in a concentration
3	ranging from about 0.02 mg/mL to about 0.5 mg/mL.
1	Claim 62 (original): A unit dosage form for the treatment of herpes simplex
2	and conditions giving rise thereto, said unit dosage form being a topical dermal dosage form
3	selected from the group consisting of topical lotions, gels, creams, and emulsions, comprising
4	as active ingredients:
5	(a) ascorbic acid,
6	(b) 2-amino-2-deoxy-D-glucose,
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7	(c) zinc sulfate, and
8	(d) L-lysine hydrochloride.
1	Claim 63 (original): A unit dosage form in accordance with claim 62 in which
2	the concentrations of said active ingredients are as follows:
3	(a) about 1.3 μg/mL to about 30 μg/mL of ascorbic acid,
4	(b) about 0.01 mg/mL to about 0.2 mg/mL of 2-amino-2-deoxy-D-glucose,
5	(c) about 0.06 μg/mL to about 8.5 μg/mL of zinc sulfate, and
6	(d) about 1.6 μ g/mL to about 23 μ g/mL of L-lysine hydrochloride.
1	Claim 64 (original): A unit dosage form in accordance with claim 63 further
2	comprising as an active ingredient Cu^{+2} in a concentration ranging from about 0.15 $\mu\text{g/mL}$ to
3	about 15 μg/mL.
1	Claim 65 (original): A unit dosage form in accordance with claim 64 further
2	comprising as an active ingredient quercetin in a concentration ranging from about 0.12
3	$\mu g/mL$ to about 2.75 $\mu g/mL$.
1	Claim 66 (original): A unit dosage form in accordance with claim 65 further
2	comprising as an active ingredient heparin sodium in a concentration ranging from about 0.6
3	unit/mL to about 8 units/mL.
1	Claim 67 (original): A unit dosage form in accordance with claim 66 further
2	comprising as an active ingredient D,α-tocopherol in a concentration ranging from about 16
3	$\mu g/mL$ to about 1600 $\mu g/mL$.
1	Claim 68 (original): A unit dosage form in accordance with claim 67 in which
2	said D,α -tocopherol is in the form of D,α -tocopherol nicotinate in a concentration ranging
3	from about 19 μg/mL to about 2600 μg/mL.

- 1 Claim 69 (original): A unit dosage form in accordance with claim 67 in which
- 2 said D,α-tocopherol is in the form of D,α-tocopherol succinate in a concentration ranging
- 3 from about 19 μ g/mL to about 2500 μ g/mL.

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